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         1 FILE CABA
         5 FILE CANCERLIT
        66 FILE CAPLUS
        85 FILE DGENE
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         2 FILE GENBANK
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        24 FILE IFIPAT
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        23 FILE MEDLINE
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       543 FILE USPATFULL
        43 FILE USPAT2
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=> (alpha with factor with leader with sequence) and (ADH2 with promoter)
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=> d ab bib 1-7

L9 ANSWER 1 OF 7 CAPLUS COPYRIGHT 2005 ACS on STN

AB Provided is a method for producing physiol. active protein in high yields by controlling the redox potential of a fermentation A physiol. active protein such as IFN-α is characteristically produced in a high yield by using Saccharomyces cerevisiae DCO4 (KCTC0051BP) which is transformed with a plasmid having ADH2/GAP promoter, alpha-factor leader sequence, and genes of IFN-alpha, adjusting pH within the range of 4 to 8 for the fermentation and metal ions as oxidants or sulfur compound as reductants to keep redox

AN 2004:856530 CAPLUS

DN 142:133164

TI Fermentation process for producing active proteins from recombinant Saccharomyces cerevisiae using redox potential control

potential value to be greater than 0 and smaller than 180.

IN Kwon, In Chan; Han, Kyu Boem

PA Lg Chemical Co., Ltd, S. Korea

SO Repub. Korea, No pp. given

CODEN: KRXXFC

DT Patent

LA Korean

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
ΡI	KR 235462	B1	19991215	KR 1997-9188	19970318
PRAI	KR 1997-9188		19970318		

AB A method for controlling oxidation and reduction of physiol. active protein, which is produced from fermentation of recombinant microorganism, is provided

by

adding oxidant and reductant. Physiol. active protein, for example, IFN- $\alpha$  is produced in a high yield by using Saccharomyces cerevisiae DCO4(KCTC 0051BP), which is transformed with a plasmid having ADH2/GAP promoter, alpha-factor leader

sequence and genes of IFN-alpha, and adding 1-50 mM of sulfur compound as a reductant into medium wherein, the sulfur compound is selected from DTT(dithiothreitol), cysteine, or  $\beta$ -mercaptoethanol.

AN 2004:856529 CAPLUS

DN 142:133163

- TI Redox potential control during Saccharomyces cerevisiae recombinant protein fermentations
- IN Kwon, In Chan; Han, Kyu Boem
- PA Lg Chemical Co., Ltd, S. Korea
- SO Repub. Korea, No pp. given CODEN: KRXXFC
- DT Patent
- LA Korean

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
ΡI	KR 235461	B1	19991215	KR 1996-39670	<b>199</b> 60913
PRAI	KR 1996-39670		19960913		

- L9 ANSWER 3 OF 7 CAPLUS COPYRIGHT 2005 ACS on STN
- Amethod for mass producing human interferon alpha(IFN-α) from the recombinant yeast is provided to improve a production yield and facilitate folding of the IFN-α. Human IFN α is mass produced from the recombinant yeast by incubating Saccharomyces cerevisiae DC04 transformed by the plasmid containing ADH2/GAP promoter, alpha -factor leader sequence, and the IFN-alpha gene in a medium containing metal ion at 25 to 30°, pH 4.5 to 5.0 under aerobic condition, adding a proper amount of glucose (avoiding accumulation in the medium), at the time of starting expression by exhaustion of glucose and ethanol. A 0.1 to 10 mM of metal ion is added into the medium to facilitate folding of the IFN-α; it contains copper, iron, zinc, manganese, molybdenum, and cobalt.
- AN 2004:852101 CAPLUS
- DN 142:79847
- TI Method for mass producing human interferon alpha(IFN- $\alpha$ ) from recombinant yeast
- IN Han, Gyu Beom; Kwon, Seon Hun
- PA Lg Chemical Co., Ltd., S. Korea
- SO Repub. Korea, No pp. given

CODEN: KRXXFC

DT Patent

LA Korean

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	KR 177321	B1	19990401	KR 1996-26844	<b>1996</b> 0703
PRAI	KR 1996-26844		19960703		

- L9 ANSWER 4 OF 7 CAPLUS COPYRIGHT 2005 ACS on STN
- AB A method for producing human granulocyte colony stimulation factor(G-CSF) from recombinant yeast is provided. Recombinant yeast Saccharomyces cerevisiae DCO4, which is transformed by a plasmid containing alc. dehydrogenase 2/glyceraldehyde-3-phosphate(ADH2/GAP) promoter, .alpha.-factor leader

sequence, and the G-CSF gene in which 17th cysteine from 5'
terminal is substituted with serine, is fed-batch fermented under
conditions of 25 to 30 °C and pH 4.5-5.0 glucose is added into

reactor in order to maintain the yeast growth rate to 0.08 to 0.12 /h when ethanol the concentration is zero.

- AN 2004:844141 CAPLUS
- DN 142:54843
- TI Human granulocyte-colony stimulating factor expression in recombinant Saccharomyces cerevisiae
- IN Han, Kyu Sum; Kwon, Sun Hoon; Lee, Heung Yeup
- PA Lg Chemical Ltd, S. Korea
- SO Repub. Korea, No pp. given CODEN: KRXXFC
- DT Patent
- LA Korean
- FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
ΡI	KR 154965	B1	19981015	KR 1996-75	19960105
PRAI	KR 1996-75		19960105		

- L9 ANSWER 5 OF 7 CAPLUS COPYRIGHT 2005 ACS on STN
- AB Transmission-blocking vaccines based on sexual-stage surface antigens of P. falciparum may assist in the control of this lethal form of human malaria. Two vaccine candidates, Pfs25 and Pfs28, were produced as single recombinant fusion proteins. The 39-kDa chimeric proteins, having a C-terminal His6 tag, were secreted by S. cerevisiae, using the prepro-alpha.-factor leader sequence.

Pfs25-28 fusion proteins were more potent than either Pfs25 or Pfs28 alone in eliciting antibodies in mice that blocked oocyst development in Anopheles freeborni mosquitoes: complete inhibition of oocyst development in the mosquito midgut was achieved with fewer vaccinations, at a lower dose, and for a longer duration than with either Pfs25 or Pfs28 alone. Increased antigen-specific IgG titers and highly significant lymphoproliferative stimulation by Pfs28-containing antigens suggest the presence of an immunodominant helper T-cell epitope in the Pfs28 portion of the fusion proteins. This epitope may be responsible for the enhanced humoral response to both Pfs25 and Pfs28 antigens. Protein production of the fusion protein was improved 12-fold by converting Pfs28 codons to yeast-preferred codons (TBV28), using a modified ADH2 promoter and incorporating a (Glu-Ala)2 repeat after the Kex2 cleavage site.

- AN 1998:7805 CAPLUS
- DN 128:113774
- TI Saccharomyces cerevisiae-secreted fusion proteins Pfs25 and Pfs28 elicit potent Plasmodium falciparum transmission-blocking antibodies in mice
- AU Gozar, Mary Margaret G.; Price, Virginia L.; Kaslow, David C.
- CS Malaria Vaccines Section, Laboratory of Parasitic Diseases, National Institutes of Health, Bethesda, MD, 20892-0425, USA
- SO Infection and Immunity (1998), 66(1), 59-64 CODEN: INFIBR; ISSN: 0019-9567
- PB American Society for Microbiology
- DT Journal
- LA English
- RE.CNT 20 THERE ARE 20 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT
- L9 ANSWER 6 OF 7 CAPLUS COPYRIGHT 2005 ACS on STN
- AB This article describes the construction of various alc. dehydrogenase 2 gene (ADH2) promoter plasmids for expression of heterologous proteins in yeast as well as for secretion into the culture medium. Plasmids YEpC-pADH2a, b, and d contain a polylinker following the ADH2 promoter. YEpC-PADH2a and YEpC-PADH2b contain the pUC13 polylinker in both orientations with unique restriction sites for SmaI, BamHI, and SalI. They do not have an ATG codon as does YEpC-PADH2d and therefore can be used for cloning and expression of intact genes. The yeast-Escherichia coli shuttle vector pαADH2 allows regulated

secretion of heterologous proteins via the ADH2 promoter fused to the .alpha.-factor leader sequence. The YIp5-derived integrating vectors pBC36 and pBC72 allow overprodn. of ADR1 which is required for pos. activation of the ADH2 promoter. The expression of several cDNAs encoding mouse and human granulocyte-macrophage colony-stimulating factor and human and bovine interleukin II was tested using vector p $\alpha$ ADH2. The amount of heterologous protein secreted ranged from 5 to 70  $\mu$ g/mL.

- AN 1990:546651 CAPLUS
- DN 113:146651
- TI Expression of heterologous proteins in Saccharomyces cerevisiae using the ADH2 promoter
- AU Price, Virginia L.; Taylor, Wayne E.; Clevenger, William; Worthington, Marlis; Young, Elton T.
- CS Dep. Mol. Biol., Immunex Corp., Seattle, WA, 98101, USA
- SO Methods in Enzymology (1990), 185(Gene Expression Technol.), 308-18 CODEN: MENZAU; ISSN: 0076-6879
- DT Journal
- LA English
- L9 ANSWER 7 OF 7 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
- AB Transmission-blocking vaccines based on sexual-stage surface antigens of Plasmodium falciparum may assist in the control of this lethal form of human malaria. Two vaccine candidates, Pfs25 and Pfs28, were produced as single recombinant fusion proteins. The 39-kDa chimeric proteins, having a C-terminal His6 tag, were secreted by Saccharomyces cerevisiae, using the prepro-alpha-factor leader

sequence. Pfs25-28 fusion proteins were significantly more potent than either Pfs25 or Pfs28 alone in eliciting antibodies in mice that blocked oocyst development in Anopheles freeborni mosquitoes: complete inhibition of oocyst development in the mosquito midgut was achieved with fewer vaccinations, at a lower dose, and for a longer duration than with either Pfs25 or Pfs28 alone. Increased antigen-specific immunoglobulin G titers and highly significant lymphoproliferative stimulation by Pfs28-containing antigens suggest the presence of an immunodominant helper T-cell epitope in the Pfs28 portion of the fusion proteins. This epitope may be responsible for the enhanced humoral response to both Pfs25 and Pfs28 antigens. Protein production of the fusion protein was improved 12-fold by converting Pfs28 codons to yeast-preferred codons (TBV28), using a modified ADH2 promoter and incorporating a (Glu-Ala)2 repeat after the Kex2 cleavage site.

- AN 1998:78406 BIOSIS
- DN PREV199800078406
- TI Saccharomyces cerevisiae-secreted fusion proteins Pfs25 and Pfs28 elicit potent Plasmodium falciparum transmission-blocking antibodies in mice.
- AU Gozar, Mary Margaret G.; Price, Virginia L.; Kaslow, David C. [Reprint author]
- CS Malaria Vaccines Section, Lab. Parasitic Diseases, Natl. Inst. Health, Build. 4, Room B1-31, Bethesda, MD 20892-0425, USA
- SO Infection and Immunity, (Jan., 1998) Vol. 66, No. 1, pp. 59-64. print. CODEN: INFIBR. ISSN: 0019-9567.
- DT Article
- LA English
- ED Entered STN: 24 Feb 1998 Last Updated on STN: 24 Feb 1998